LISTING OF THE CLAIMS

- 1. (Withdrawn) A composition comprising erythropoietin and an erythropoietin production inducing peptide (EPIP).
- 2. (Previously Presented) A composition comprising an erythropoietin production inducing peptide (EPIP), wherein the EPIP comprises poly-D-glutamic acid, poly-L-glutamic acid, poly-D-aspartic acid, poly-L-aspartic acid, or a mixture of both.
- 3. (Withdrawn) The composition of claim 103, comprising a therapeutically effective amount of erythropoietin and EPIP.
- 4. (Withdrawn) The composition of claim 2, comprising a therapeutically effective amount of an EPIP.
- 5. (Withdrawn Currently Amended) The composition of <u>claim 104</u> elaim 103, wherein the erythropoietin is recombinant erythropoietin.
- 6-14. (Canceled)
- 15. (Previously Presented) The composition of claim 2, wherein the EPIP is poly-D-glutamic acid.
- 16-18. (Canceled)
- 19. (Previously Presented) The composition of claim 2, further comprising a pharmaceutically acceptable diluent, adjuvant or carrier.
- 20. (Previously Presented) The composition of claim 2, wherein the preservative comprises benzyl alcohol, a paraben and phenol, or a mixture thereof.
- 21. (Previously Presented) The composition of claim 2, wherein the composition further comprises a buffering agent.
- 22. (Original) The composition of claim 21, wherein the buffering agent comprises citrate, phosphate, tartrate, succinate, adipate, maleate, lactate and acetate buffers, sodium bicarbonate, and sodium carbonate, or a mixture thereof.
- 23. (Previously Presented) The composition of claim 2, further comprising an isotonicity adjusting agent, wherein the isotonicity adjusting agent comprises sodium chloride, glycerol, mannitol, sorbitol, or a mixture thereof.

- 24. (Previously Presented) The composition of claim 2, further comprising a pH adjusting agent that adjusts the pH of the solution within the range of 5-8.
- 25. (Previously Presented) The composition of claim 2, further comprising human serum albumin.
- 26. (Previously Presented) The composition of claim 2, wherein the composition is an aqueous solution, a non-aqueous suspension, or a dry powder.
- 27. (Previously Presented) The composition of claim 2, wherein the composition is in oral dosage form.
- 28. (Previously presented) The composition of claim 2, further comprising fatty acid(s), surfactant(s), or enteric material, or a mixture thereof, wherein components are mixed in liquid phase and lyophilized.
- 29. (Previously Presented) The composition of claim 2, wherein the composition is in injectable form.
- 30. (Withdrawn) A composition comprising cells derived either from a subject treated with EPIP or co-cultured *in vitro* with proximal tubular cells exposed to EPIP either *in vivo* or *ex vivo*, wherein the derived cells can be propagated *in vitro* and are capable of producing erythropoietin, wherein said EPIP is poly-D-glutamic acid, poly-L- glutamic acid, poly-D-aspartic acid, poly-L-aspartic acid, or a mixture of both.
- 31. (Canceled)
- 32. (Withdrawn) The composition of claim 30, wherein said EPIP is poly-D-glutamic acid.
- 33. (Withdrawn) The composition of claim 30, wherein said cells produce 100 U or more of erythropoietin per 10⁶ cells in 48 hours.
- 34. (Withdrawn) The composition of claim 32, wherein the cells are capable of producing 500 U or more of erythropoietin.
- 35. (Withdrawn) The composition of claim 32, wherein the cells are capable of producing 1000 U or more of erythropoietin.
- 36. (Withdrawn) The composition of claim 30, wherein the poly-D-glutamic acid is administered in the amount of about 50 mg/kg of body weight per day to 400 mg/kg of body weight per day.

- 37. (Withdrawn) The composition of claim 30, wherein the cells continue to express erythropoietin after exposure to the EPIP.
- 38. (Withdrawn) The composition of claim 30, wherein the cells are indirectly stimulated to produce erythropoietin by the EPIP.
- 39. (Withdrawn) The composition of claim 30, wherein proximal tubular cells are treated with EPIP and in turn stimulate proliferation of erythropoietin producing cells.
- 40. (Withdrawn) A method of treatment comprising administering erythropoietin to a subject, wherein the erythropoietin is produced by the method of claim 74.
- 41. (Withdrawn) A method of treatment comprising administering erythropoietin and an EPIP to a subject, wherein the EPIP comprises poly-D-glutamic acid, poly-L-glutamic acid, poly-D-aspartic acid, poly-L-aspartic acid, or a mixture of both.
- 42-51. (Canceled)
- 52. (Withdrawn) The method of claim 41, wherein the EPIP comprises poly-D-glutamic acid.
- 53-55. (Canceled)
- 56. (Withdrawn) The method of claim 41, wherein the method of treatment comprises treating anemia, Crohn's Disease, ulcerative colitis, chronic renal insufficiency, or end stage renal disease or any erythropoietin-responsive anemia.
- 57. (Withdrawn) The method of claim 41, wherein the treatment results in angiogenesis in the kidney.
- 58. (Withdrawn) The method of claim 41, wherein the method of treatment comprises treating organ or tissue transplantation subjects.
- 59. (Withdrawn) The method of claim 41, wherein the method of treatment comprises enhancing wound healing.
- 60. (Canceled)
- 61. (Withdrawn) The method of claim 41, wherein the subject is a mammal.
- 62. (Withdrawn) The method of claim 41, wherein the subject is human.
- 63. (Withdrawn) The method of claim 41, wherein the erythropoietin and/or EPIP is administered by intravenous or intramuscular or subcutaneous or intraperitoneal injection.

- 64. (Withdrawn) The method of any one of claims 63, wherein the erythropoietin, EPIP, or erythropoietin and EPIP is administered orally or rectally.
- 65. (Withdrawn) The method of claim 41, wherein a mechanical device directs a stream of a therapeutically effective amount of poly-D-glutamic acid into the oral cavity of a mammal while the mammal is inhaling.
- 66. (Withdrawn) The method of claim 65, wherein the mechanical device is selected from the group consisting of a nebulizer, a metered dose inhaler, and a powder inhaler.
- 67. (Withdrawn) The method of claim 41, wherein the administration of poly-D-glutamic acid results in a red blood cell level of 5000 or more erythrocytes per μL of blood.
- 68-71. (Canceled)
- 72. (Withdrawn Currently Amended) A method for the production of erythropoietin, the method comprising: a) contacting cells in culture with a EPIP, wherein the EPIP comprises poly-D-glutamic acid, poly-L-glutamic acid, poly-D-aspartic acid, poly-L-aspartic acid, or a mixture of both, and b) harvesting erythropoietin from these cells.
- 73. (Withdrawn Currently Amended) A method for the production of erythropoietin, comprising a) administering a EPIP to a mammal, wherein the EPIP comprises poly-D-glutamic acid, poly-L-glutamic acid, poly-D-aspartic acid, poly-L-aspartic acid, or a mixture of both, and b) harvesting erythropoietin producing cells from the mammal.
- 74. (Withdrawn Currently Amended) A method for the production of erythropoietin comprising a) administering a EPIP to a mammal, wherein the EPIP comprises poly-D-glutamic acid, poly-L-glutamic acid, poly-D-aspartic acid, poly-L-aspartic acid, or a mixture of both, and b) harvesting proximal tubular cells from the kidney.
- 75. (Withdrawn) The method of claim 74, wherein said EPIP comprises poly-D- glutamic acid.
- 76. (Withdrawn) The method of claim 74, wherein the erythropoietin is harvested by steps comprising: a) removing culture fluid from the cells; and b) isolating erythropoietin from the culture fluid.
- 77. (Withdrawn) The method of claim 76, wherein the erythropoietin is isolated from the culture fluid by using HPLC.

- 78. (Withdrawn) The method of claim 74, wherein erythropoietin is not isolated from the cell culture.
- 79. (Canceled)
- 80. (Canceled)
- 81. (Withdrawn) The method of claim 79, wherein the cells are peritubular insterstitial cells.
- 82. (Withdrawn) The method of claim 80, wherein co-cultures of proximal tubular cells cause the proliferation of fibroblast cells.
- 83. (Withdrawn) The method of claim 79, wherein the cells are kidney cells.
- 84. (Withdrawn) The method of claim 74, wherein at least 50% of the cells are producing erythropoietin.
- 85-97. (Canceled)
- 98. (Withdrawn Currently Amended) A method of making cells that produce erythropoietin comprising administering to the cells an effective amount of an EPIP, wherein the EPIP comprises poly-D-glutamic acid, poly-L-glutamic acid, poly-D-aspartic acid, poly-L-aspartic acid, or a mixture of both.
- 99. (Withdrawn) The method of claim 98, wherein the EPIP is administered to a cell that then stimulates the erythropoietin producing cell to produce erythropoietin.
- 100. (Withdrawn) The method of claim 99, wherein the cell that stimulates the erythropoietin producing cell is a proximal tubular cell.
- 101. (Withdrawn) The method of claim 98, wherein the cells continue to produce erythropoietin after exposure to EPIP.
- 102. (Canceled)
- 103. (Withdrawn) The composition of claim 2, wherein the composition further comprises erythropoietin.
- 104. (New) The composition of claim 2, wherein the composition further comprises erythropoietin.